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            IN THE UNITED STATES DISTRICT COURT FOR THE
2
                   NORTHERN DISTRICT OF OKLAHOMA
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4
     W. A. DREW EDMONDSON, in his )
5
     capacity as ATTORNEY GENERAL )
     OF THE STATE OF OKLAHOMA and )
6
     OKLAHOMA SECRETARY OF THE
     ENVIRONMENT C. MILES TOLBERT,)
7
     in his capacity as the
     TRUSTEE FOR NATURAL RESOURCES)
8
     FOR THE STATE OF OKLAHOMA,
9
                  Plaintiff,
10
                                    )4:05-CV-00329-TCK-SAJ
     VS.
11
     TYSON FOODS, INC., et al,
12
                  Defendants.
13
14
                       THE VIDEOTAPED DEPOSITION OF
15
     VALERIE HARDWOOD, PhD, produced as a witness on
16
     behalf of the Defendants in the above styled and
17
     numbered cause, taken on the 18th day of July, 2008,
18
     in the City of Tulsa, County of Tulsa, State of
19
     Oklahoma, before me, Lisa A. Steinmeyer, a Certified
20
     Shorthand Reporter, duly certified under and by
21
     virtue of the laws of the State of Oklahoma.
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1	A P P E A R A N C E S
2	HOD WILL DI ATAMETERS
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5	-and- Mr. Louis Bullock
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TULSA FREELANCE REPORTERS 918-587-2878

I N D E X WITNESS P A G E VALERIE HARWOOD Direct Examination by Mr. Todd Direct Examination by Ms. Longwell Signature Page Reporter's Certificate

1	A	Of course, I've done some additional data		
2	analysis for the report.			
3	Q	Right, and you submitted a report?		
4	A	Correct.		
5	Q	We talked at your last deposition you	09:09AM	
6	talke	d at your last deposition a bit about fate and		
7	transı	port, and let me just run through some		
8	chara	cteristics here, and I hope we can take care of		
9	these	pretty quickly. Since your prior deposition,		
10	have y	you conducted any study of the fate and	09:09AM	
11	transport characteristics of any bacterium in the			
12	Illinois River watershed?			
13	A	No, I have not.		
14	Q	So you have not studied how bacteria is		
15	affected by temperature? 09:09AM		09:09AM	
16	A	No.		
17	Q	Desiccation?		
18	A	No.		
19	Q	Predation?		
20	A	No.	09:09AM	
21	Q	Osmotic pressure?		
22	A	No.		
23	Q	UV exposure?		
24	A	No.		
25	Q	pH balance?	09:09AM	

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1	A	No.	
2	Q	Nutrient availability?	
3	A	No.	
4	Q	Have you studied how the movement of any	
5	partic	cular bacterium in the IRW is affected by its	09:09AM
6	size?		
7	A	No, I have not.	
8	Q	Its shape?	
9	A	No.	
10	Q	It's surface charge?	09:10AM
11	A	No.	
12	Q	Location in the water column?	
13	A	No.	
14	Q	Presence of vegetation?	
15	A	No.	09:10AM
16	Q	The media it's moving through?	
17	A	No.	
18	Q	Have you cultured the Brevibacterium that you	
19	identi	ified through your PCR process?	
20	A	No.	09:10AM
21	Q	Why not?	
22	A	There has been no need to culture the	
23	Brevik	pacterium.	
24	Q	Have you identified it any more specifically	
25	than t	to say it's 98 percent consistent with	09:10AM

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1	Brevibacteria avium?		
2	A No.		
3	Q And if you haven't cultured, I assume	e you also	
4	have not studied its fate and transport		
5	characteristics?	09:10AM	
6	A That's correct.		
7	Q Now, what you refer to as the marker	, the	
8	biomarker in your term, what you're actually	Y	
9	referring to is actually the DNA sequence the	nat's	
10	contained by the Brevibacterium; is that con	rrect? 09:10AM	
11	A That is correct. We're referring to	the DNA	
12	sequence, yes.		
13	Q Okay. For clarity, I'm going to atte	empt to be	
14	consistent referring to the Brevibacterium	as the	
15	PCR Brevibacterium and the sequence as the	PCR 09:10AM	
16	sequence. Will those terms make sense to you? I		
17	just want to distinguish the two.		
18	A Well, it's really a DNA sequence, so	I	
19	guess		
20	Q We can call it the DNA sequence.	09:11AM	
21	A DNA sequence.		
22	Q If I refer to that, then we're talking	ng about	
23	what you would refer to as the biomarker?		
24	A Yes.		
25	Q Now, we previously discussed or at ye	our last 09:11AM	

1	deposition you discussed that when a bacteria dies,		
2	its DNA remains in the environment for some period		
3	of time after that. Do you recall that?		
4	A Yes, it can remain for some period of time.		
5	Q Do you know how long the DNA sequence at issue 09:11AM		
6	in this case can remain in nature apart from the		
7	Brevibacterium that carries it?		
8	A Typically in nature, bacterial DNA is rapidly		
9	degraded within and it depends on the		
10	environment, but within a matter of hours to several 09:11AM		
11	days.		
12	Q Okay. You said it depends on the environment.		
13	A Correct.		
14	Q What kind of characteristics affect how		
15	quickly the DNA degrades? 09:11AM		
16	A Characteristics would include the amount of		
17	ultraviolet radiation. It would include the amount		
18	of pred or not predation but the amount of		
19	organisms that would consume that DNA because		
20	they'll use it as a food source. So it would depend 09:12AM		
21	on the trophic level. So in a more eutrophic		
22	nutrient dense environment, then that DNA would		
23	probably be consumed more quickly than in a more		
24	allegatory thick environment.		
25	Q Can DNA move in the environment after the 09:12AM		

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1	bacte	ria that carried it had died, become inactive?	
2	A	DNA could be transported along with water,	
3	yes.		
4	Q	Could it move in any other way?	
5	A	It would not be able to be motile on its own.	09:12AM
6	So it	would have to be transported by the movement	
7	of wat	ter or some other matrix.	
8	Q	Okay. Let's talk briefly about sources of	
9	bacte	ria in the IRW. Since your last deposition,	
10	have y	you studied sources in the IRW, apart from	09:13AM
11	poulti	ry, of any of fecal indicator bacteria?	
12	A	I have not.	
13	Q	Okay. Has anyone associated with the State's	
14	case?		
15	A	Roger Olsen of CDM has done some work with	09:13AM
16	bacte	ria in cow manure.	
17	Q	Okay. Are you familiar with the nature of his	
18	work?		
19	A	I have read his report, yes.	
20	Q	Have you studied any sources in the IRW, apart	09:13AM
21	from poultry, of E. coli?		
22	A	No, I have not.	
23	Q	Okay. Of Enterococci?	
24	A	No, I have not.	
25	Q	Campylobacter?	09:13AM

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1	A	No.		
2	Q	Salmonella?		
3	A	No.		
4	Q	Any other bacteria?		
5	A	No.	09:13AM	
6	Q	Have you undertaken yourself to quantify fecal		
7	produc	ction levels by any animal in the IRW?		
8	A	No, I have not.		
9	Q	Have you undertaken quantification of bacteria		
10	loadir	ng from any particular source in the IRW?	09:13AM	
11	A	I have not.		
12	Q	Now, you submitted a journal article to the		
13	Journa	al of Applied and Environmental Microbiology;		
14	correct?			
15	A	That's correct.	09:14AM	
16	Q	And we were provided a copy of that a couple		
17	of day	ys ago. You're on the editorial board of that		
18	journa	al?		
19	A	That's correct.		
20	Q	Okay. Have you discussed your article with	09:14AM	
21	any of	your colleagues on that board?		
22	A	No, I have not. That wouldn't be you don't		
23	do tha	at.		
24	Q	Okay. You submitted it on June 11, at least		
25	accord	ding to the cover E-mail; is that correct?	09:14AM	

1	contamination.	
2	Q Okay, but in order for it to be an indicator	
3	of poultry fecal contamination, is it necessary that	
4	the PCR sequence share the same fate and transport	
5	as pathogens from poultry litter?	02:00PM
6	A Can you say that again? I just got to get the	
7	first part.	
8	Q Sure. In order for it to be an indicator	
9	you've just said it is an	
10	A Indicator of poultry fecal contamination.	02:00PM
11	Q Right, and that fecal contamination you are	
12	talking about here is bacteria; correct?	
13	A Correct.	
14	Q Okay. So in order for the presence of the	
15	indicator	02:00PM
16	A I'm sorry. Let me go back there because we're	
17	not only concerned about bacterial fecal	
18	contamination from poultry, we're also concerned	
19	about nutrient contamination. So we can add	
20	nutrients and metals to that list.	02:00PM
21	Q We'll talk about let's table the nutrients	
22	and the metals for just a second and let's talk	
23	about bacteria. In order for it to indicate the	
24	presence of bacteria derived from poultry, is it	
25	necessary that the PCR that the Brevibacterium	02:00PM

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1	that you identified share the fate and transport	
2	characteristics of other bacteria from poultry	
3	litter?	
4	A It would have to have certain fate and	
5	transport characteristics in common. 02:01PM	
6	Q Okay. If we compare the correlations that we	
7	discussed here, so the correlation, let's say,	
8	taking Enterococcus, for instance, the relationship	
9	between Enterococcus and the sequence in litter as	
10	.75 and the relationship between Enterococcus and 02:01PM	
11	the biomarker the sequence in water is .89, which	
12	is different; correct?	
13	A It's different, but it's certainly within the	
14	bounds of what you would expect from regular	
15	sampling error. 02:01PM	
16	Q Okay. How big a difference can you have	
17	within the bounds of regular sampling error?	
18	A In environmental microbiology we're very happy	
19	to get correlations of .3 as long as they're	
20	statistically significant, even .2 sometimes. So 02:01PM	
21	there's a really wide range of what you can get from	
22	correlations and still be biologically meaningful.	
23	Q Okay. So does it surprise you at all then	
24	that the correlation that you got between E. coli	
25	and the PCR sequence in litter was .39 you told me 02:02PM	